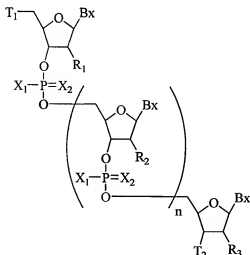


This listing of claims will replace all prior versions, and listings, of claims in the application.

**Listing of Claims:**

1. (currently amended) An oligomeric compound having the formula:



wherein:

each Bx is, independently, a heterocyclic base moiety;

T<sub>2</sub> is hydroxyl[[,] or a protected hydroxyl,~~an oligonucleotide or an oligonucleoside;~~

T<sub>1</sub> is a modified phosphate having the formula:



wherein

Q is OH or CH<sub>3</sub>

R<sub>1</sub>, R<sub>3</sub> and each R<sub>2</sub> are, independently, hydrogen, hydroxyl, a sugar substituent group or a protected sugar substituent group;

each X<sub>1</sub> and X<sub>2</sub> is, independently, O or S wherein at least one X<sub>1</sub> is S; and  
 n is from 3 to 48.

- 2-3. (canceled)
4. (previously presented) The oligomeric compound of claim 1 wherein Q is CH<sub>3</sub>.
- 5-10. (canceled)
11. (original) The oligomeric compound of claim 1 wherein R<sub>1</sub>, R<sub>3</sub> and each R<sub>2</sub> is hydrogen.
12. (original) The oligomeric compound of claim 1 wherein R<sub>1</sub>, R<sub>3</sub> and each R<sub>2</sub> is hydroxyl.
13. (previously presented) The oligomeric compound of claim 1 wherein R<sub>1</sub>, R<sub>3</sub> and each R<sub>2</sub> are, independently, hydrogen, hydroxyl, a sugar substituent group or a protected sugar substituent group.
14. (original) The oligomeric compound of claim 1 wherein at least one of R<sub>1</sub>, R<sub>2</sub> or R<sub>3</sub> is an optionally protected sugar substituent group.
15. (original) The oligomeric compound of claim 1 wherein each X<sub>2</sub> is S.
16. (original) The oligomeric compound of claim 1 wherein each heterocyclic base moiety is, independently, adenine, cytosine, 5-methylcytosine, thymine, uracil, guanine or 2-aminoadenine.
17. (original) The oligomeric compound of claim 1 wherein n is from about 8 to about 30.
18. (original) The oligomeric compound of claim 1 wherein n is from about 15 to 25.

19. (withdrawn) A method of treating an organism having a disease characterized by the undesired production of a protein comprising contacting the organism with an oligomeric compound of claim 1.

20. (previously presented) A composition comprising:  
a pharmaceutically effective amount of an oligomeric compound of claim 1; and  
a pharmaceutically acceptable diluent or carrier.

21. (withdrawn) A method of modifying *in vitro* a nucleic acid, comprising contacting a test solution containing RNase H and said nucleic acid with an oligomeric compound of claim 1.

22. (withdrawn) A method of concurrently enhancing hybridization and RNase H activation in a organism comprising contacting the organism with an oligomeric compound of claim 1.

23. (withdrawn) A method comprising contacting a cell with an oligomeric compound of claim 1.

24-41. (canceled)